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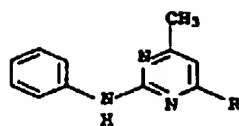
(74) **ROBIC**

(54) **MELANGES FONGICIDES A BASE DE COMPOSES AMIDE ET DE DERIVES PYRIDINE**
(54) **FUNGICIDE MIXTURES BASED ON PYRIDINE
CARBOXAMIDES**

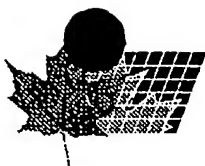
(57)

The invention relates to fungicide mixtures containing, as active components, a) an amide compound of formula (I) A-CO-NR¹R², wherein A represents an aryl group or an aromatic or non-aromatic, 5- or 6-structured heterocyclic compound having 1 to 3 heteroatoms selected from O, N, S; whereby the aryl group or the heterocyclic compound can optionally comprise 1, 2 or 3 substituents selected independently of one another from alkyl, halogen, CHF₂, CF₃, alkoxyl, haloalkoxyl, alkylthio, alkyl sulfinyl and alkyl sulfonyl; R¹ represents a hydrogen atom; R² represents a phenyl group or cycloalkyl group optionally containing 1, 2 or 3 substituents selected independently of one another from alkyl, alkenyl, alkynyl, alkoxyl, alkenyloxy, alkynyloxy, cycloalkyl, cycloalkenyl, cycloalkyloxy, cycloalkenyloxy, phenyl and halogen, whereby the aliphatic and cycloaliphatic radicals can be partially or completely halogenated and/or the cycloaliphatic radicals can be substituted by 1 to 3 alkyl groups, whereby the phenyl group can contain 1 to 5 halogen atoms and/or 1 to 3 substituents selected independently of one another from alkyl, haloalkyl, alkoxyl, haloxy, alkylthio and haloalkylthio, and whereby the amidic phenyl group is optionally condensed with a saturated 5-structured ring which is optionally substituted by one or more alkyl groups and/or can comprise a heteroatom selected from O and S, and b) fungicides from the group of dicarboximides, and/or c) pyrimidine derivatives of formula (III) in which R represents methyl, propyne-1-yl or cyclopropyl, and/or d) fludioxinil or fenpiclonil and/or captan, captafol or folpet, and/or f)

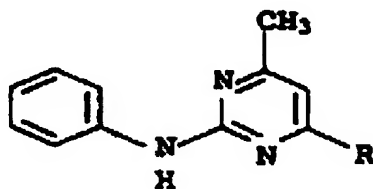
fluazinam, and/or g) dichlofuanid or tolyl fuanid.
The active components are provided in a
synergistically effective quantity.



(III)



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(30) 1997/12/18 (19756379.1) DE
(54) **MELANGES FONGICIDES A BASE DE COMPOSES AMIDE ET
DE DERIVES PYRIDINE**
(54) **FUNGICIDE MIXTURES BASED ON PYRIDINE
CARBOXAMIDES**

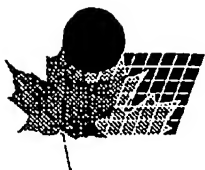


(III)

(57) L'invention concerne des mélanges fongicides contenant une quantité à action synergique des constituants actifs suivants: a) un composé amide de la formule (I) $\Lambda\text{-CO-NR}^1\text{R}^2$, dans laquelle Λ désigne un groupe aryle ou un hétérocycle aromatique ou non aromatique ayant 5 ou 6 chaînons, qui peut comporter entre 1 et 3 hétéroatomes, sélectionnés parmi O, N et S; le groupe aryle ou l'hétérocycle pouvant éventuellement comporter 1, 2 ou 3 substituants, sélectionnés indépendamment les uns des autres parmi alkyle, halogène, CHF_2 , CF_3 , alcoxy, halogénure d'alcoxy, alkylthio, alkylsulfinyle et alkylsulfonyl; R^1 désigne un

(57) The invention relates to fungicide mixtures containing, as active components, a) an amide compound of formula (I) $\Lambda\text{-CO-NR}^1\text{R}^2$, wherein Λ represents an aryl group or an aromatic or non-aromatic, 5- or 6-structured heterocyclic compound having 1 to 3 heteroatoms selected from O, N, S; whereby the aryl group or the heterocyclic compound can optionally comprise 1, 2 or 3 substituents selected independently of one another from alkyl, halogen, CHF_2 , CF_3 , alkoxyl, haloalkoxyl, alkylthio, alkyl sulfinyl and alkyl sulfonyl; R^1 represents a hydrogen atom; R^2 represents a phenyl group or cycloalkyl group optionally containing 1, 2 or 3





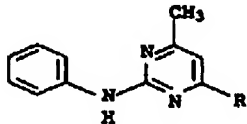
(21) (A1) **2,313,322**
(86) 1998/12/15
(87) 1999/07/01

atome d'hydrogène; R^2 désigne un groupe phényle ou cycloalkyle, présentant éventuellement 1, 2 ou 3 substituants, sélectionnés indépendamment les uns des autres parmi alkyle, alkényle, alkynyle, alcoxy, alkényloxy, alkynyloxy, cycloalkyle, cycloalkényle, cycloalkyloxy, cycloalkényloxy, phényle et halogène, les restes aliphatiques et cycloaliphatiques pouvant être partiellement ou entièrement halogénés et/ou les restes cycloaliphatiques pouvant être substitués par 1 à 3 groupes alkyle et le groupe phényle pouvant comporter entre 1 et 5 atomes d'halogène et/ou entre 1 et 3 substituants, sélectionnés indépendamment les uns des autres, parmi alkyle, halogénure d'alkyle, alcoxy, halogénure d'alcoxy, alkylthio et halogénure d'alkylthio, le groupe phényle amidique étant éventuellement condensé avec un composé cyclique saturé à 5 chaînons, qui est éventuellement substitué par un ou plusieurs groupes alkyle, et/ou un hétéroatome, sélectionné parmi O et S, et b) des fongicides du groupe des dicarboximides et/ou c) des dérivés de pyrimidine de la formule (III), dans laquelle R désigne méthyle, propine-1-yle ou cyclopropyle, et/ou d) fludioxinil ou fenpiclonil et/ou e) captane, captafol ou folpet et/ou f) fluazinam et/ou g) dichlofluanide ou tolylfluanide.

substituents selected independently of one another from alkyl, alkenyl, alkynyl, alkoxy, alkenyloxy, alkynyloxy, cycloalkyl, cycloalkenyl, cycloalkyloxy, cycloalkenyloxy, phenyl and halogen, whereby the aliphatic and cycloaliphatic radicals can be partially or completely halogenated and/or the cycloaliphatic radicals can be substituted by 1 to 3 alkyl groups, whereby the phenyl group can contain 1 to 5 halogen atoms and/or 1 to 3 substituents selected independently of one another from alkyl, haloalkyl, alkoxy, haloxy, alkylthio and haloalkylthio, and whereby the amidic phenyl group is optionally condensed with a saturated 5-structured ring which is optionally substituted by one or more alkyl groups and/or can comprise a heteroatom selected from O and S, and b) fungicides from the group of dicarboximides, and/or c) pyrimidine derivatives of formula (III) in which R represents methyl, propyne-1-yl or cyclopropyl, and/or d) fludioxinil or fenpiclonil and/or captan, captafol or folpet, and/or f) fluazinam, and/or g) dichlofluanid or tolyl fluanid. The active components are provided in a synergistically effective quantity.

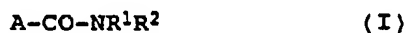


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INTERNATIONALE ZUSAMMENARBEIT AUF DEM GEBIET DES PATENTWESENS (PCT)

(51) Internationale Patentklassifikation ⁶ : A01N 43/00	A2	(11) Internationale Veröffentlichungsnummer: WO 99/31981 (43) Internationales Veröffentlichungsdatum: 1. Juli 1999 (01.07.99)
(21) Internationales Aktenzeichen: PCT/EP98/08231 (22) Internationales Anmeldedatum: 15. Dezember 1998 (15.12.98) (30) Prioritätsdaten: 197 56 379.1 18. Dezember 1997 (18.12.97) DE (71) Anmelder (für alle Bestimmungsstaaten ausser US): BASF AKTIENGESELLSCHAFT [DE/DE]; D-67056 Ludwigshafen (DE). (72) Erfinder; und (75) Erfinder/Anmelder (nur für US): SCHELBERGER, Klaus [AT/DE]; Traminerweg 2, D-67161 Gönnsheim (DE). SCHERER, Maria [DE/DE]; Hermann-Jürgens-Strasse 30, D-76829 Landau (DE). EICKEN, Karl [DE/DE]; Am Hüttenwingert 12, D-67157 Wachenheim (DE). HAMPEL, Manfred [DE/DE]; Im Blengarten 15, D-67435 Neustadt (DE). AMMERMAN, Eberhard [DE/DE]; Von-Gagem-Strasse 2, D-64646 Heppenheim (DE). LORENZ, Gisela [DE/DE]; Erlenweg 13, D-67434 Neustadt (DE). STRATHMANN, Siegfried [DE/DE]; Donnersbergstrasse 9, D-67117 Limburgerhof (DE).	(74) Gemeinsamer Vertreter: BASF AKTIENGESELLSCHAFT; D-67056 Ludwigshafen (DE). (81) Bestimmungsstaaten: AL, AU, BG, BR, BY, CA, CN, CZ, GE, HU, ID, IL, JP, KR, KZ, LT, LV, MK, MX, NO, NZ, PL, RO, RU, SG, SI, SK, TR, UA, US, eurasisches Patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), europäisches Patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Veröffentlicht <i>Ohne internationalen Recherchenbericht und erneut zu veröffentlichen nach Erhalt des Berichts.</i>	
(54) Title: FUNGICIDE MIXTURES BASED ON AMIDE COMPOUNDS AND PYRIDINE DERIVATIVES		
(54) Bezeichnung: FUNGIZIDE MISCHUNGEN AUF DER BASIS VON AMIDVERBINDUNGEN UND PYRIDINDERIVATEN		
<div style="text-align: center;">  </div> <div style="text-align: right;">(II)</div>		
(57) Abstract <p>The invention relates to fungicide mixtures containing, as active components, a) an amide compound of formula (I) $A-CO-NR^1R^2$, wherein A represents an aryl group or an aromatic or non-aromatic, 5- or 6-structured heterocyclic compound having 1 to 3 heteroatoms selected from O, N, S; whereby the aryl group or the heterocyclic compound can optionally comprise 1, 2 or 3 substituents selected independently of one another from alkyl, halogen, CHF_2, CF_3, alkoxy, haloalkoxy, alkylthio, alkyl sulfinyl and alkyl sulfonyl; R^1 represents a hydrogen atom; R^2 represents a phenyl group or cycloalkyl group optionally containing 1, 2 or 3 substituents selected independently of one another from alkyl, alkenyl, alkynyl, alkoxy, alkenyloxy, alkynyloxy, cycloalkyl, cycloalkenyl, cycloalkyloxy, cycloalkenyloxy, phenyl and halogen, whereby the aliphatic and cycloaliphatic radicals can be partially or completely halogenated and/or the cycloaliphatic radicals can be substituted by 1 to 3 alkyl groups, whereby the phenyl group can contain 1 to 5 halogen atoms and/or 1 to 3 substituents selected independently of one another from alkyl, haloalkyl, alkoxy, haloxy, alkylthio and haloalkylthio, and whereby the amidic phenyl group is optionally condensed with a saturated 5-structured ring which is optionally substituted by one or more alkyl groups and/or can comprise a heteroatom selected from O and S, and b) fungicides from the group of dicarboximides, and/or c) pyrimidine derivatives of formula (III) in which R represents methyl, propyne-1-yl or cyclopropyl, and/or d) fludioxinil or fenpiclonil and/or captan, captafol or folpet, and/or f) fluazinam, and/or g) dichlofuanid or tolyl fuanid. The active components are provided in a synergistically effective quantity.</p>		

FUNGICIDE MIXTURES BASED ON PYRIDINE CARBOXAMIDES

The present invention relates to fungicidal mixtures for controlling harmful fungi, which mixtures comprise amide compounds of the formula I



in which

- 10 A is an aryl group or an aromatic or non-aromatic, 5- or 6-membered heterocycle which has from 1 to 3 hetero atoms selected from O, N and S;
- where the aryl group or the heterocycle may or may not have 1, 2 or 3 substituents which are selected, independently of one another, from alkyl, halogen, CHF_2 , CF_3 , alkoxy, haloalkoxy, alkylthio, alkylsulfynyl and alkylsulfonyl;
- R^1 is a hydrogen atom;
- R^2 is a phenyl or cycloalkyl group which may or may not have 1, 2 or 3 substituents which are selected from alkyl, alkenyl, alkynyl, alkoxy, alkenyloxy, alkynyloxy, cycloalkyl, cycloalkenyl, cycloalkyloxy, cycloalkenyloxy, phenyl and halogen, where the aliphatic and cycloaliphatic radicals may be partially or fully halogenated and/or the cycloaliphatic radicals may be substituted by from 1 to 3 alkyl groups and
- 20 where the phenyl group may have from 1 to 5 halogen atoms and/or from 1 to 3 substituents which are selected, independently of one another, from alkyl, haloalkyl, alkoxy, haloalkoxy, alkylthio and haloalkylthio, and where the amidic phenyl group may or may not be condensed with a saturated 5-membered ring which may or may not be substituted by one or more alkyl groups and/or may have a hetero atom selected from O and S,

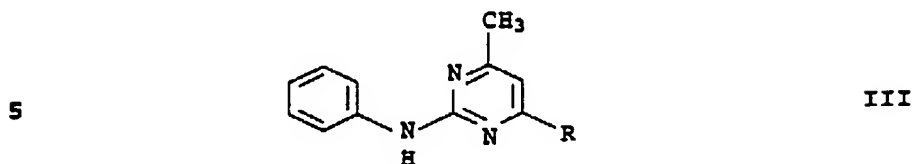
and

b) fungicides of the group of the dicarboximides

30 and/or

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c) a pyrimidine derivative of the formula III,



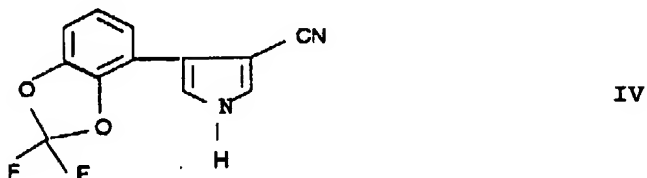
in which R is methyl, propyn-1-yl or cyclopropyl,

10 and/or

d) at least one active ingredient of the formula IV or V,

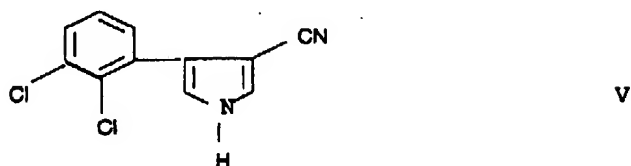
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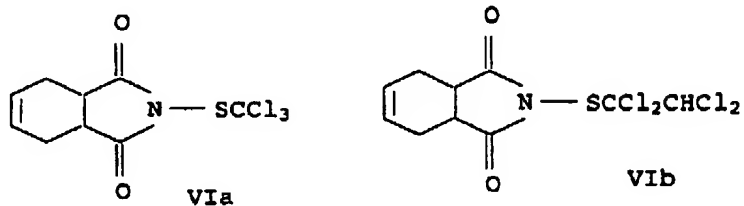


35 and/or

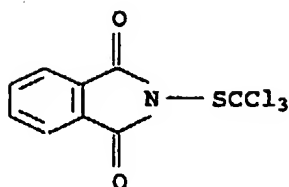
e) a phthalimide derivative selected from the group consisting of the compounds VIa, VIb and VII

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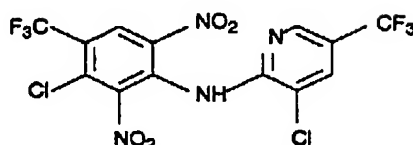


VII

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and/or

- 10 f) a dinitroaniline of the formula VIII



VIII

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and/or

- 20 g) an arylsulfamide of the formula IX a or IXb



IXa

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CH₃

IXb

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in a synergistically effective amount.

WO 97/08952 describes fungicidal mixtures which, in addition to compounds of the formula I, also comprise fenazaquin as further components. These are described as being very effective against Botrytis.

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The amide compounds of the formula I are known per se and are described in the literature (EP-A 545 099).

- 40 Fungicides from the group of the dicarboximides are known to the person skilled in the art and are commercially available.

Also known are the pyrimidine derivatives III, their preparation and their activity against harmful fungi [R = methyl: DD-A 151 404 (common name: pyrimethanil); R = 1-propynyl: EP-A 224 339 (common name: mepanipyrin); R=cyclopropyl: EP-A 310 550].

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The compound IV is known from K. Gehmann, R. Nyfeler, A.J. Leadbeater, D. Nevill and D. Sozzi, Proceedings of the Brighton Crop Protection Conference, Pests and Diseases 1990, Vol. 2, p. 399 (common name: fludioxinil) and is commercially available from Novartis.

10 The compound V is known from D. Nevill, R. Nyfeler, D. Sozzi, Proceedings of the Brighton Crop Protection Conference, Pests and Diseases 1988, Vol. 1, p. 65 (common name: fenpiclonil).

US-A 2,553,770; 2,553,771; 2,553,776 describe the compounds VIa (common name: captan) and VII (common name: folpet), their preparation and their activity against harmful fungi. The compound VIb (common name: captafol) is described in
15 Phytopathology 52(1962), 52, 754.

Likewise, the compound VIII, its preparation and its use (CAS RN 79622-59-6, common name: fluazinam) are described in the
20 literature.

The compounds IXa and IXb are known under the common names dichlofluanid and tolylfluanid, respectively, and are described in the German patent 1193498.
25

It is an object of the present inventions [sic] to provide mixtures which have an improved activity against harmful fungi combined with a reduced total amount of active ingredients applied (synergistic mixtures), with a view to reducing the application rates and to improving the activity spectrum of the
30 known compounds.

We have found that this object is achieved by the mixtures defined at the outset. Moreover, we have found that better
35 control of harmful fungi is possible by applying the compounds I and the compounds II to VIII simultaneously, that is either together or separately, or by applying the compounds I and the compounds II to VIII in succession than when the compounds I or II to VIII are applied on their own.
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The mixtures according to the invention have synergistic action and are therefore particularly suitable for controlling harmful fungi, in particular of Botrytis species.

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In the context of the present invention, halogen is fluorine, chlorine, bromine and iodine and is in particular fluorine, chlorine and bromine.

- 5 The term "alkyl" includes straight-chain and branched alkyl groups. These are preferably straight-chain or branched C₁-C₁₂-alkyl and in particular C₁-C₆-alkyl groups. Examples of alkyl groups are alkyl such as, in particular, methyl, ethyl, propyl, 1-methylethyl, butyl, 1-methylpropyl, 2-methylpropyl, 10 1,1-dimethylethyl, n-pentyl, 1-methylbutyl, 2-methylbutyl, 3-methylbutyl, 1,2-dimethylpropyl, 1,1-dimethylpropyl, 2,2-dimethylpropyl, 1-ethylpropyl, n-hexyl, 1-methylpentyl, 2-methylpentyl, 3-methylpentyl, 4-methylpentyl, 1,2-dimethylbutyl, 1,3-dimethylbutyl, 2,3-dimethylbutyl, 15 1,1-dimethylbutyl, 2,2-dimethylbutyl, 3,3-dimethylbutyl, 1,1,2-trimethylpropyl, 1,2,2-trimethylpropyl, 1-ethylbutyl, 2-ethylbutyl, 1-ethyl-2-methylpropyl, n-heptyl, 1-methylhexyl, 1-ethylpentyl, 2-ethylpentyl, 1-propylbutyl, octyl, decyl, dodecyl.

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- Haloalkyl is an alkyl group as defined above which is partially or fully halogenated by one or more halogen atoms, in particular by fluorine and chlorine. Preferably, there are from 1 to 3 halogen atoms present, and the difluoromethane [sic] or the 25 trifluoromethyl group is particularly preferred.

- The above statements for the alkyl group and the haloalkyl group apply in a corresponding manner to the alkyl and haloalkyl groups 30 in alkoxy, haloalkoxy, alkylthio, haloalkylthio, alkylsulfynyl and alkylsulfonyl.

- The alkenyl group includes straight-chain and branched alkenyl groups. These are preferably straight-chain or branched 35 C₃-C₁₂-alkenyl groups and in particular C₃-C₆-alkenyl groups. Examples of alkenyl groups are 2-propenyl, 2-butenyl, 3-butenyl, 1-methyl-2-propenyl, 2-methyl-2-propenyl, 2-pentenyl, 3-pentenyl, 4-pentenyl, 1-methyl-2-butenyl, 2-methyl-2-butenyl, 3-methyl-2-butenyl, 1-methyl-3-butenyl, 2-methyl-3-butenyl, 40 3-methyl-3-butenyl, 1,1-dimethyl-2-propenyl, 1,2-dimethyl-2-propenyl, 1-ethyl-2-propenyl, 2-hexenyl, 3-hexenyl, 4-hexenyl, 5-hexenyl, 1-methyl-2-pentenyl, 2-methyl-2-pentenyl, 3-methyl-2-pentenyl, 4-methyl-2-pentenyl, 1-methyl-3-pentenyl, 2-methyl-3-pentenyl, 3-methyl-3-pentenyl, 45 4-methyl-3-pentenyl, 1-methyl-4-pentenyl, 2-methyl-4-pentenyl, 3-methyl-4-pentenyl, 4-methyl-4-pentenyl, 1,1-dimethyl-2-butenyl, 1,1-dimethyl-3-butenyl, 1,1-dimethyl-3-butenyl [sic],

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1,2-dimethyl-2-butenyl, 1,2-dimethyl-3-butenyl,
 1,3-dimethyl-2-butenyl, 1,3-dimethyl-3-butenyl,
 2,2-dimethyl-3-butenyl, 2,3-dimethyl-2-butenyl,
 2,3-dimethyl-3-butenyl, 1-ethyl-2-butenyl, 1-ethyl-3-butenyl,
 5 2-ethyl-2-butenyl, 2-ethyl-3-butenyl, 1,1,2-trimethyl-2-propenyl,
 1-ethyl-1-methyl-2-propenyl and 1-ethyl-2-methyl-2-propenyl, in
 particular 2-propenyl, 2-butenyl, 3-methyl-2-butenyl and
 3-methyl-2-pentenyl.

10 The alkenyl group may be partially or fully halogenated by one or
 more halogen atoms, in particular by fluorine or chlorine. The
 alkenyl group preferably has from 1 to 3 halogen atoms.

15 The alkynyl group includes straight-chain and branched alkynyl
 groups. These are preferably straight-chain and branched
 C₃-C₁₂-alkynyl groups and in particular C₃-C₆-alkynyl groups.
 Examples of alkynyl groups are 2-propynyl, 2-butyne, 3-butyne,
 1-methyl-2-propynyl, 2-pentyne, 3-pentyne, 4-pentyne,
 20 1-methyl-3-butyne, 2-methyl-3-butyne, 1-methyl-2-butyne,
 1,1-dimethyl-2-propynyl, 1-ethyl-2-propynyl, 2-hexynyl,
 3-hexynyl, 4-alkynyl [sic], 5-hexynyl, 1-methyl-2-pentyne,
 1-methyl-3-pentyne, 1-methyl-4-pentyne, 2-methyl-3-pentyne,
 2-methyl-4-pentyne, 3-methyl-4-pentyne, 4-methyl-2-pentyne,
 1,2-dimethyl-2-butyne, 1,1-dimethyl-3-butyne,
 25 1,2-dimethyl-3-butyne, 2,2-dimethyl-3-butyne,
 1-ethyl-2-butyne, 1-ethyl-3-butyne, 2-ethyl-3-butyne and
 1-ethyl-1-methyl-2-propynyl.

30 The above statements for the alkenyl group and its halogen
 substituents and for the alkynyl group apply in a corresponding
 manner to alkenyloxy and alkynyloxy.

The cycloalkyl group is preferably a C₃-C₆-cycloalkyl group, such
 35 as cyclopropyl, cyclobutyl, cyclopentyl or cyclohexyl. If the
 cycloalkyl group is substituted, it preferably has from 1 to 3
 C₁-C₄-alkyl radicals as substituents.

Cycloalkenyl is preferably a C₄-C₆-cycloalkenyl group, such as
 40 cyclobutenyl, cyclopentenyl or cyclohexenyl. If the cycloalkenyl
 group is substituted, it preferably has from 1 to 3 C₁-C₄-alkyl
 radicals as substituents.

7

A cycloalkoxy group is preferably a C₅-C₆-cycloalkoxy group, such as cyclopentyloxy or cyclohexyloxy. If the cycloalkoxy group is substituted, it preferably has from 1 to 3 C₁-C₄-alkyl radicals as substituents.

5

The cycloalkenyloxy group is preferably a C₃-C₆-cycloalkenyloxy group, such as cyclopentyloxy or cyclohexyloxy. If the cycloalkenyloxy group is substituted, it preferably has from 1 to 3 C₁-C₄-alkyl radicals as substituents.

10

Aryl is preferably phenyl.

If A is a phenyl group, this may have one, two or three of the abovementioned substituents in any position. These substituents are preferably selected, independently of one another, from alkyl, difluoromethyl, trifluoromethyl and halogen, in particular chlorine, bromine and iodine. Particularly preferably, the phenyl group has a substituent in the 2-position.

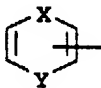
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If A is a 5-membered heterocycle, it is in particular a furyl, thiazolyl, pyrazolyl, imidazolyl, oxazolyl, thienyl, triazolyl or thiadiazolyl radical or the corresponding dihydro or tetrahydro derivatives thereof. Preference is given to a thiazolyl or pyrazolyl radical.

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If A is a 6-membered heterocycle, it is in particular a pyridyl radical or a radical of the formula:

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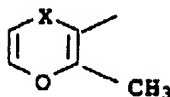
in which one of the radicals X and Y is O, S or NR²³, where R²³ is H or alkyl, and the other of the radicals X and Y is CH₂, S, SO, SO₂ or NR²³. The dotted line means that a double bond may or may not be present.

40

The 6-membered aromatic heterocycle is particularly preferably a pyridyl radical, in particular a 3-pyridyl radical, or a radical of the formula

45

8



(A3)

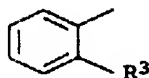
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10 in which X is CH₂, S, SO or SO₂.

The abovementioned heterocyclic radicals may or may not have 1, 2 or 3 of the abovementioned substituents, where these substituents are preferably selected, independently of one another, from
15 alkyl, halogen, difluoromethyl or trifluoromethyl.

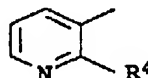
A is particularly preferably a radical of the formulae:

20

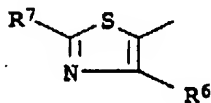


(A1)

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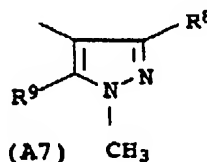


(A2)



(A5)

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(A7)

in which R³, R⁴, R⁶, R⁷, R⁸ and R⁹ independently of one another are
35 hydrogen, alkyl, in particular methyl, halogen, in particular chlorine, CHF₂ or CF₃.

The radical R¹ in the formula I is preferably a hydrogen atom.

40

The radical R² in the formula I is preferably a phenyl radical. R² preferably has at least one substituent which is particularly preferably in the 2-position. The substituent (or the substituents) is (are) preferably selected from the group
45 consisting of alkyl, cycloalkyl, cycloalkenyl, halogen or phenyl.

The substituents of the radical R^2 may in turn be substituted again. The aliphatic or cycloaliphatic substituents may be partially or fully halogenated, in particular fluorinated or chlorinated. They preferably have 1, 2 or 3 fluorine or chlorine atoms. If the substituent of the radical R^2 is a phenyl group, this phenyl group may preferably be substituted by from 1 to 3 halogen atoms, in particular chlorine atoms, and/or by a radical which is preferably selected from alkyl and alkoxy. Particularly preferably, the phenyl group is substituted with a halogen atom in the p-position, i.e. the particularly preferred substituent of the radical R^2 is a p-halogen-substituted phenyl radical. The radical R^2 may also be condensed with a saturated 5-membered ring, where this ring for its part may have from 1 to 3 alkyl substituents.

15

R^2 is in this case, for example, indanyl, thiaindanyl and oxaindanyl. Preference is given to indanyl and 2-oxaindanyl which are attached to the nitrogen atom in particular via the 4-position.

20

According to a preferred embodiment, the composition according to the invention comprises as amide compound a compound of the formula I in which A is as defined below:

25

phenyl, pyridyl, dihydropyranyl, dihydrooxathiynyl, dihydrooxathiynyloxide, dihydrooxathiynyldioxide, furyl, thiazolyl, pyrazolyl or oxazolyl, where these groups may have 1, 2 or 3 substituents which are selected, independently of one another, from alkyl, halogen, difluoromethyl and trifluoromethyl.

30

According to a further preferred embodiment, A is one of the following groups:

pyridin-3-yl, which may or may not be substituted in the 2-position by halogen, methyl, difluoromethyl, trifluoromethyl, methoxy, methylthio, methylsulfinyl or methylsulfonyl;

35

phenyl, which may or may not be substituted in the 2-position by methyl, trifluoromethyl, chlorine, bromine or iodine;

40

2-methyl-5,6-dihydropyran-3-yl;

2-methyl-5,6-dihydro-1,4-oxathiyn-3-yl or the 4-oxide or 4,4-dioxide thereof;

45

10

2-methylfuran-3-yl, which may or may not be substituted in the 4- and/or 5-position by methyl;

5 thiazol-5-yl, which may or may not be substituted in the 2- and/or 4-position by methyl, chlorine, difluoromethyl or trifluoromethyl;

10 thiazol-4-yl, which may or may not be substituted in the 2- and/or 5-position by methyl, chlorine, difluoromethyl or trifluoromethyl;

15 1-methylpyrazol-4-yl, which may or may not be substituted in the 3- and/or 5-position by methyl, chlorine, difluoromethyl or trifluoromethyl; or

oxazol-5-yl, which may or may not be substituted in the 2- and/or 4-position by methyl or chlorine.

20 According to a further preferred embodiment, the compositions according to the invention comprise as amide compound a compound of the formula I in which R² is a phenyl group which may or may not be substituted by 1, 2 or 3 of the abovementioned substituents.

25 According to a further preferred embodiment, the compositions according to the invention comprise as amide compound a compound of the formula I in which R² is a phenyl group which has one of the following substituents in the 2-position:

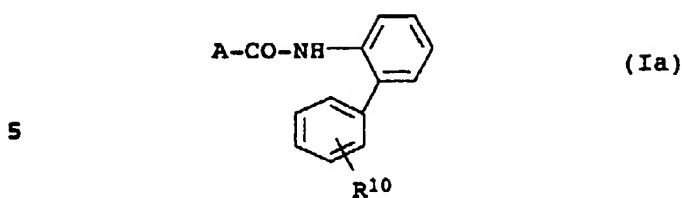
30 C₃-C₆-alkyl, C₅-C₆-cycloalkenyl, C₅-C₆-cycloalkyloxy, cycloalkenyloxy, where these groups may be substituted by 1, 2 or 3 C₁-C₄-alkyl groups,

35 phenyl, which is substituted by from 1 to 5 halogen atoms and/or from 1 to 3 groups which are selected, independently of one another, from C₁-C₄-alkyl, C₁-C₄-haloalkyl, C₁-C₄-alkoxy, C₁-C₄-haloalkoxy, C₁-C₄-alkylthio and C₁-C₄-haloalkylthio,

40 indanyl or oxaindanyl which may or may not be substituted by 1, 2 or 3 C₁-C₄-alkyl groups.

45 According to a further preferred embodiment, the compositions according to the invention comprise as amide compound a compound of the formula Ia,

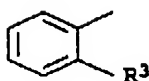
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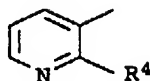
10 in which

A is

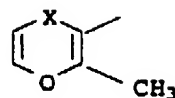
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(A1)



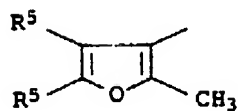
(A2)



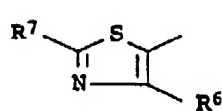
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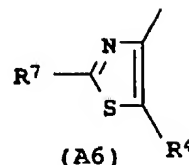
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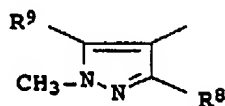


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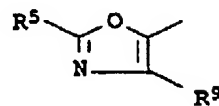


(A6)

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(A7)



(A8)

35

X is methylene, sulfur, sulfinyl or sulfonyl (SO₂),

R³ is methyl, difluoromethyl, trifluoromethyl, chlorine, bromine or iodine,

40 R⁴ is trifluoromethyl or chlorine,

R⁵ is hydrogen or methyl,

R⁶ is methyl, difluoromethyl, trifluoromethyl or chlorine,

R⁷ is hydrogen, methyl or chlorine,

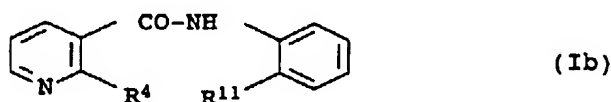
45 R⁸ is methyl, difluoromethyl or trifluoromethyl,

12

R⁹ is hydrogen, methyl, difluoromethyl, trifluoromethyl or chlorine,
R¹⁰ is C₁-C₄-alkyl, C₁-C₄-alkoxy, C₁-C₄-alkylthio or halogen.

5 According to a particularly preferred embodiment, the compositions comprise as amide compound a compound of the formula Ib

10



15

in which

R⁴ is halogen and

R¹¹ is phenyl which is substituted by halogen.

20

Useful amide compounds of the formula I are mentioned in EP-A-545 099 and 589 301 which are incorporated herein in their entirety by reference.

25

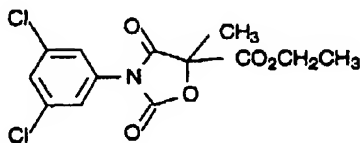
The preparation of the amide compounds of the formula I is known, for example, from EP-A-545 099 or 589 301 or can be carried out by similar processes.

30 Preferred substituents R²¹ and R²² are hydrogen, F, Cl, methyl, ethyl, methoxy, thiomethyl and N-methylamino. R³ and R⁴ together may also form a grouping =O.

35 Preferred fungicides of the dicarboximide type are the compounds IIa to IIId:

II.a: ethyl (")-3-(3,5-dichlorophenyl)-5-methyl-2,4-dioxoxazolidine-5-carboxylate [sic]

40



45

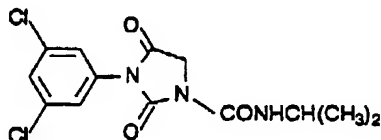
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II.b: 3-(3,5-dichlorophenyl)-N-isopropyl-2,4-dioxoimidazolidine-1-carboxamide

5



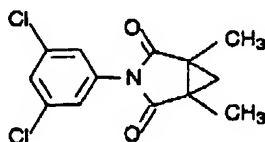
II.b

or

10

II.c: N-(3,5-dichlorophenyl)-1,2-dimethylcyclopropane-1,2-dicarboximide

15



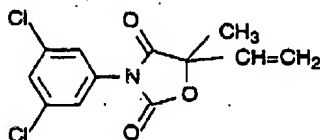
II.c

or

20

II.d: (RS)-3-(3,5-dichlorophenyl)-5-ethenyl-5-methyl-1,3-oxazolidine-2,4-dione

25



II.d.

These compounds are commercially available as fungicidally active ingredients and are described in the literature as follows:

- II.a (common name: chlozolate): CAS RN [84332-86-5], DE-A 29 06 574;
- 35 • II.b (common name: iprodione): CAS RN [36734-19-7], US-A 3,755,350;
- II.c (common name: procymidone): CAS RN [32809-16-8], US-A 3,903,090;
- 40 • II.d (common name: vinclozoline): CAS RN [50471-44-8], DE-A 22 07 576].

To unfold the synergistic activity, even a small amount of the amide compound of the formula I is sufficient. Preference is given to employing amide compound and active ingredient of the formula II and/or III to IX in a weight ratio in the range of
45 from 50:1 to 1:50, in particular from 10:1 to 1:10. It is also possible here to employ ternary mixtures which, in addition to

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amide compounds I, comprise both compounds II and one or more compounds III to IX. In such mixtures, the mixing ratio of the compounds II and III to IX with each other is usually in the range of from 50:1 to 1:50, preferably from 10:1 to 1:10.

5

Owing to the basic character of their nitrogen atoms, the compounds III, IV, V and VIII are capable of forming salts or adducts with inorganic or organic acids or with metal ions.

10

Examples of inorganic acids are hydrohalic acids such as hydrofluoric acid, hydrochloric acid, hydrobromic acid and hydriodic acid, sulfuric acid, phosphoric acid and nitric acid.

15

Suitable organic acids are, for example, formic acid, carbonic acid [sic] and alkanolic acids, such as acetic acid, trifluoroacetic acid, trichloroacetic acid and propionic acid, and also glycolic acid, thiocyanic acid, lactic acid, succinic acid, citric acid, benzoic acid, cinnamic acid, oxalic acid,

20

alkylsulfonic acids (sulfonic acids having straight-chain or branched alkyl radicals of 1 to 20 carbon atoms), arylsulfonic acids or aryldisulfonic acids (aromatic radicals, such as phenyl and naphthyl, which carry one or two sulfo groups), alkylphosphonic acids (phosphonic acids having straight-chain or

25

branched alkyl radicals of 1 to 20 carbon atoms), arylphosphonic acids or aryldiphosphonic acids (aromatic radicals, such as phenyl and naphthyl, which carry one or two phosphoric [sic] acid radicals), it being possible for the alkyl or aryl radicals to carry further substituents, eg. p-toluenesulfonic acid, salicylic

30

acid, p-aminosalicylic acid, 2-phenoxybenzoic acid, 2-acetoxybenzoic acid, etc.

Suitable metal ions are, in particular, the ions of the elements of the first to eighth sub-group, in particular chromium,

35

manganese, iron, cobalt, nickel, copper, zinc and furthermore of the second main group, in particular calcium and magnesium, and of the third and fourth main group, in particular aluminum, tin and lead. The metals can exist in the various valencies which they can assume.

40

When preparing the mixtures, it is preferred to employ the pure active ingredients I and II to IX, to which further active ingredients against harmful fungi or other pests, such as insects, arachnids or nematodes, or else herbicidal or

45

growth-regulating active ingredients or fertilizers, can be admixed.

15

The mixtures of the compounds I and II and/or III to IX, or the compounds I and II and/or III to IX used simultaneously, jointly or separately, exhibit outstanding activity against a wide range of phytopathogenic fungi, in particular from the classes of the
5 Ascomycetes, Basidiomycetes, Phycomycetes and Deuteromycetes. Some of them act systemically and can therefore be employed as foliar- and soil-acting fungicides.

They are especially important for controlling a large number of
10 fungi in a variety of crop plants, such as cotton, vegetable species (eg. cucumbers, beans, tomatoes, potatoes and cucurbits), barley, grass, oats, bananas, coffee, maize, fruit species, rice, rye, soya, grapevine, wheat, ornamentals, sugar cane, and a variety of seeds.

15

They are particularly suitable for controlling the following phytopathogenic fungi: Erysiphe graminis (powdery mildew) in cereals, Erysiphe cichoracearum and Sphaerotheca fuliginea in cucurbits, Podosphaera leucotricha in apples, Uncinula necator in
20 grapevines, Puccinia species in cereals, Rhizoctonia species in cotton, rice and lawns, Ustilago species in cereals and sugar cane, Venturia inaequalis (scab) in apples, Helminthosporium species in cereals, Septoria nodorum in wheat, Botrytis cinerea [sic] (gray mold) in strawberries, vegetables, ornamentals and
25 grapevines, Cercospora arachidicola in groundnuts, Pseudocercospora herpotrichoides in wheat and barley, Pyricularia oryzae in rice, Phytophthora infestans in potatoes and tomatoes, Plasmopara viticola in grapevines, Pseudoperonospora species in hops and cucumbers, Alternaria
30 species in vegetables and fruit, Mycosphaerella species in bananas and Fusarium and Verticillium species.

The mixtures according to the invention may particularly
35 preferably be employed for controlling Botrytis species in crops of grapevines and vegetables, and also in ornamentals.

The compounds I and II and/or III to IX can be applied simultaneously, either together or separately, or in succession,
40 the sequence, in the case of separate application, generally not having any effect on the result of the control measures.

Depending on the kind of effect desired, the application rates of the mixtures according to the invention are, in particular in
45 agricultural crop areas, from 0.01 to 8 kg/ha, preferably 0.1 to 5 kg/ha, in particular 0.2 to 3.0 kg/ha.

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The application rates of the compounds I are from 0.01 to 2.5 kg/ha, preferably 0.05 to 2.5 kg/ha, in particular 0.1 to 1.0 kg/ha.

- 5 Correspondingly, in the case of the compounds II and/or III to IX, the application rates are from 0.01 to 10 kg/ha, preferably 0.05 to 5 kg/ha, in particular 0.05 to 2.0 kg/ha.
- 10 For seed treatment, the application rates of the mixture are generally from 0.001 to 250 g/kg of seed, preferably 0.01 to 100 g/kg, in particular 0.01 to 50 g/kg.

- If phytopathogenic harmful fungi are to be controlled, the
- 15 separate or joint application of the compounds I and II and/or III to IX or of the mixtures of the compounds I and II and/or III to IX is effected by spraying or dusting the seeds, the plants or the soils before or after sowing of the plants, or before or after plant emergence.

- 20 The fungicidal synergistic mixtures according to the invention, or the compounds I and II and/or III to IX, can be formulated for example in the form of ready-to-spray solutions, powders and suspensions or in the form of highly concentrated aqueous, oily
- 25 or other suspensions, dispersions, emulsions, oil dispersions, pastes, dusts, materials for broadcasting or granules, and applied by spraying, atomizing, dusting, broadcasting or watering. The use form depends on the intended purpose; in any case, it should ensure as fine and uniform as possible a
- 30 distribution of the mixture according to the invention.

- The formulations are prepared in a known manner, eg. by extending the active ingredient with solvents and/or carriers, if desired using emulsifiers and dispersants, it being possible also to use
- 35 other organic solvents as auxiliary solvents if water is used as the diluent. Suitable auxiliaries for this purpose are essentially: solvents such as aromatics (eg. xylene), chlorinated aromatics (eg. chlorobenzenes), paraffins (eg. mineral oil fractions), alcohols (eg. methanol, butanol), ketones (eg.
- 40 cyclohexanone), amines (eg. ethanolamine, dimethylformamide) and water; carriers such as ground natural minerals (eg. kaolins, clays, talc, chalk) and ground synthetic minerals (eg. finely divided silica, silicates); emulsifiers such as nonionic and anionic emulsifiers (eg. polyoxyethylene fatty alcohol ethers,
- 45 alkylsulfonates and arylsulfonates) and dispersants such as lignosulfite waste liquors and methylcellulose.

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Suitable surfactants are the alkali metal salts, alkaline earth metal salts and ammonium salts of aromatic sulfonic acids, eg. ligno-, phenol-, naphthalene- and dibutylnaphthalenesulfonic acid, and of fatty acids, alkyl- and alkylarylsulfonates, alkyl, 5 lauryl ether and fatty alcohol sulfates, and salts of sulfated hexa-, hepta- and octadecanols, or of fatty alcohol glycol ethers, condensates of sulfonated naphthalene and its derivatives with formaldehyde, condensates of naphthalene or of the naphthalenesulfonic acids with phenol and formaldehyde, 10 polyoxyethylene octylphenol ether, ethoxylated isooctyl-, octyl- or nonylphenol, alkylphenol polyglycol ethers, tributylphenyl polyglycol ethers, alkylaryl polyether alcohols, isotridecyl alcohol, fatty alcohol/ethylene oxide condensates, ethoxylated castor oil, polyoxyethylene alkyl ethers or polyoxypropylene 15 [sic], lauryl alcohol polyglycol ether acetate, sorbitol esters, lignosulfite waste liquors or methylcellulose.

Powders, materials for broadcasting and dusts can be prepared by mixing or jointly grinding the compounds I or II and/or III to 20 IX, or the mixture of the compounds I and II and/or III to IX, with a solid carrier.

Granules (eg. coated granules, impregnated granules or 25 homogeneous granules) are usually prepared by binding the active ingredient, or active ingredients, to a solid carrier.

Fillers or solid carriers are, for example, mineral earths, such as silica gel, silicas, silica gels [sic], silicates, talc, 30 kaolin, limestone, lime, chalk, bole, loess, clay, dolomite, diatomaceous earth, calcium sulfate, magnesium sulfate, magnesium oxide, ground synthetic materials and fertilizers, such as ammonium sulfate, ammonium phosphate, ammonium nitrate, ureas, and products of vegetable origin, such as cereal meal, tree bark 35 meal, wood meal and nutshell meal, cellulose powders or other solid carriers.

The formulations generally comprise from 0.1 to 95% by weight, preferably 0.5 to 90% by weight, of one of the compounds I and II 40 and/or III to IX or of the mixture of the compounds I and II and/or III to IX. The active ingredients are employed in a purity of from 90% to 100%, preferably 95% to 100% (according to NMR or HPLC spectrum [sic]).

45 The compounds I and II and/or III to IX, the mixtures, or the corresponding formulations, are applied by treating the harmful fungi, their habitat, or the plants, seeds, soils, areas,

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materials or spaces to be kept free from them with a fungicidally effective amount of the mixture, or of the compounds I and II and/or III to IX in the case of separate application.

- 5 Application can be effected before or after infection by the harmful fungi.

Examples of such preparations comprising the active ingredients
10 are:

- I. A solution of 90 parts by weight of the active ingredients and 10 parts by weight of N-methylpyrrolidone; this solution is suitable for use in the form of microdrops;
- 15 II. A mixture of 20 parts by weight of the active ingredients, 80 parts by weight of xylene, 10 parts by weight of the adduct of 8 to 10 mol of ethylene oxide to 1 mol of oleic acid N-monoethanolamide, 5 parts by weight of the calcium salt of dodecylbenzenesulfonate, 5 parts by weight of the
20 adduct of 40 mol of ethylene oxide and 1 mol of castor oil; a dispersion is obtained by finely distributing the solution in water;
- III. An aqueous dispersion of 20 parts by weight of the active ingredients, 40 parts by weight of cyclohexanone, 30 parts
25 by weight of isobutanol, 20 parts by weight of the adduct of 40 mol of ethylene oxide and 1 mol of castor oil;
- IV. An aqueous dispersion of 20 parts by weight of the active ingredients, 25 parts by weight of cyclohexanol, 65 parts
30 by weight of a mineral oil fraction of boiling point 210 to 280°C, and 10 parts by weight of the adduct of 40 mol of ethylene oxide and 1 mol of castor oil;
- V. A mixture, ground in a hammer mill, of 80 parts by weight of the active ingredients, 3 parts by weight of the sodium salt of diisobutyl naphthalene-1-sulfonate, 10 parts by
35 weight of the sodium salt of a lignosulfonic acid from a sulfite waste liquor and 7 parts by weight of pulverulent silica gel; a spray mixture is obtained by finely distributing the mixture in water;
- 40 VI. An intimate mixture of 3 parts by weight of the active ingredients and 97 parts by weight of finely divided kaolin; this dust comprises 3% by weight of active ingredient;

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- VII. An intimate mixture of 30 parts by weight of the active ingredients, 92 parts by weight of pulverulent silica gel and 8 parts by weight of paraffin oil which had been sprayed onto the surface of this silica gel; this formulation imparts good adhesion to the active ingredient;
- VIII. A stable aqueous dispersion of 40 parts by weight of the active ingredients, 10 parts by weight of the sodium salt of a phenolsulfonic acid/urea/formaldehyde condensate, 2 parts by weight of silica gel and 48 parts by weight of water; this dispersion may be diluted further;
- IX. A stable oily dispersion of 20 parts by weight of the active ingredients, 2 parts by weight of the calcium salt of dodecylbenzenesulfonate, 8 parts by weight of fatty alcohol polyglycol ether, 20 parts by weight of the sodium salt of a phenolsulfonic acid/urea/formaldehyde condensate and 88 parts by weight of a paraffinic mineral oil.

Use Example

- The synergistic activity of the mixtures according to the invention can be demonstrated by the following experiments:

The active ingredients, separately or together, are formulated as a 10% emulsion in a mixture of 63% by weight of cyclohexanone and 27% by weight of emulsifier, and diluted with water to the desired concentration.

- Evaluation is carried out by determining the infected leaf areas in percent. These percentages are converted into efficacies. The efficacy (W) is calculated as follows using Abbot's formula:

$$W = (1 - \alpha) \cdot 100 / \beta$$

- α corresponds to the fungal infection of the treated plants in % and
- β corresponds to the fungal infection of the untreated (control) plants in %

An efficacy of 0 means that the infection level of the treated plants corresponds to that of the untreated control plants; an efficacy of 100 means that the treated plants were not infected.

20

The expected efficacies of the mixtures of the active ingredients were determined using Colby's formula [R.S. Colby, Weeds 15, 20-22 (1967)] and compared with the observed efficacies.

5

Colby formula: $E = x + y - x \cdot y / 100$

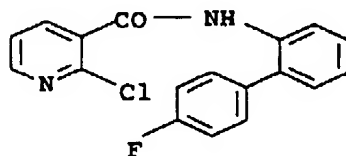
- E expected efficacy, expressed in % of the untreated control, when using the mixture of the active ingredients A and B at the concentrations a and b
- 10 x efficacy, expressed in % of the untreated control, when using active ingredient A at a concentration of a
- y efficacy, expressed in % of the untreated control, when using active ingredient B at a concentration of b
- 15

Use Example 1 - Activity against Botrytis cinerea on bell pepper leaves

- 20 Bell pepper seedlings of the variety "Neusiedler Ideal Elite" were, after 4-5 leaves were well developed, sprayed to runoff point with an aqueous preparation of active ingredient which had been prepared from a stock solution comprising 10% of active ingredient, 63% of cyclohexanone and 27% of emulsifier. The next day, the treated plants were inoculated with a spore suspension
- 25 of Botrytis cinerea containing 1.7×10^6 spores per ml of a 2% strength aqueous Biomalz solution. The test plants were subsequently placed in a climatized chamber with high atmospheric humidity at 22-24°C. After 5 days, the extent of the fungal infection on the leaves could be determined visually in %.
- 30

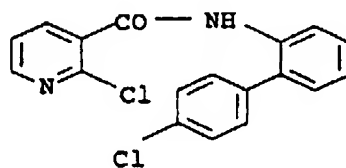
The compounds of the formula I used were the following components:

35



I.1

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I.2

45

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The results are shown in Tables 1 and 2 below.

Table 1

5	Ex.	Active ingredient	Concentration of active ingredient in the spray liquor in ppm	Efficacy in % of the untreated control
	1C	untreated	0 (99% infection)	0
10	2C	Compound I.1	31 16 8	78 67 11
	3C	Compound I.2	31 16 8	67 44 44
15	4C	Compound VIa	31 16 8	0 0 0
	5C	Compound VII	31 16 8	0 0 0
20	6C	Compound IXa	31 16 8	0 0 0

25 Table 2

	Ex.	Mixtures according to the invention (content in ppm)	observed efficacy	calculated efficacy*)
30	7	31 ppm I.1 + 31 ppm VIa	97	78
	8	8 ppm I.1 + 8 ppm VIa	30	11
35	9	31 ppm I.2 + 31 ppm VIa	100	67
	10	16 ppm I.2 + 16 ppm VIa	70	44
40	11	31 ppm I.1 + 31 ppm VII	90	78
45	12	8 ppm I.1 + 8 ppm VII	39	11

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Ex.	Mixtures according to the invention (content in ppm)	observed efficacy	calculated efficacy*)
5	13 31 ppm I.2 + 31 ppm VII	93	67
	14 16 ppm I.2 + 16 ppm VII	90	44
10	15 8 ppm I.2 + 8 ppm VII	90	44
	16 31 ppm I.1 + 31 ppm IXa	95	78
15	17 16 ppm I.1 + 16 ppm IXa	90	67
	18 31 ppm I.2 + 31 ppm IXa	100	67
20	19 16 ppm I.2 + 16 ppm IXa	99	44
25	20 8 ppm I.2 + 8 ppm IXa	90	44

*) calculated using Colby's formula

30 The test results show that for all mixing ratios the observed efficacy is higher than the efficacy which had been calculated beforehand using Colby's formula.

Use Example 2 - Activity against Botrytis cinerea on bell peppers

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40 Disks of green bell peppers were sprayed to runoff point with an aqueous preparation of active ingredient which had been prepared from a stock solution comprising 10% of active ingredient, 63 & [sic] of cyclohexanone and 27% of emulsifier. 2 hours after the spray coating had dried on, the fruit disks were inoculated with a spore suspension of Botrytis cinerea containing 1.7×10^6 spores per ml of a 2% strength Biomalz solution. The disks incubated [sic]. The Botrytis infection on the diseased fruit disks was then evaluated visually.

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The results are shown in Tables 1 [sic] and 2 [sic] below.

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Table 3

5	Ex.	Active ingredient	Concentration of active ingredient in the spray liquor in ppm	Efficacy in % of the untreated control
	21C	untreated	0 (100% infection)	0
	22C	Compound I.1	31 16	0 0
10	23C	Compound I.2	31 16	20 0
	24C	Compound IV	31 16	0 0
	25C	Compound VIa	31	0
15	26C	Compound VII	31	0
	27C	Compound VIII	31 16	0 0

20 Table 4

	Ex.	Mixtures according to the invention (content in ppm)	observed efficacy	calculated efficacy*)
25	28	31 ppm I.1 + 31 ppm IV	50	0
	29	16 ppm I.1 + 16 ppm IV	20	0
30	30	31 ppm I.2 + 31 ppm IV	70	20
	31	16 ppm I.2 + 16 ppm IV	20	0
35	32	31 ppm I.2 + 31 ppm IVa	50	20
	33	31 ppm I.2 + 31 ppm VII	40	20
40	34	31 ppm I.1 + 31 ppm VIII	30	0
45	35	16 ppm I.2 + 16 ppm VIII	20	0

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*) calculated using Colby's formula

The test results show that for all mixing ratios the observed efficacy is higher than the efficacy which had been calculated 5 beforehand using Colby's formula.

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We claim:

5 1. A fungicidal mixture, comprising as active components

a) an amide compound of the formula I



I

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in which

15 A is an aryl group or an aromatic or non-aromatic, 5- or 6-membered heterocycle which has from 1 to 3 hetero atoms selected from O, N and S;

where the aryl group or the heterocycle may or may not have 1, 2 or 3 substituents which are selected, independently of one another, from alkyl, halogen, CHF_2 , CF_3 , alkoxy, haloalkoxy, alkylthio, alkylsulfynyl and alkylsulfonyl;

20

R^1 is a hydrogen atom;

25 R^2 is a phenyl or cycloalkyl group which may or may not have 1, 2 or 3 substituents which are selected, independently of one another, from alkyl, alkenyl, alkynyl, alkoxy, alkenyloxy, alkynyloxy, cycloalkyl, cycloalkenyl, cycloalkyloxy, cycloalkenyloxy, phenyl and halogen, where the aliphatic and cycloaliphatic radicals may be partially or fully halogenated and/or the cycloaliphatic radicals may be substituted by from 1 to 3 alkyl groups and where the phenyl group may have from 1 to 5 halogen atoms and/or from 1 to 3 substituents which are selected, independently of one another, from alkyl, haloalkyl, alkoxy, haloalkoxy, alkylthio and haloalkylthio, and where the amidic phenyl group may or may not be condensed with a saturated 5-membered ring which may or may not be substituted by one or more alkyl groups and/or may have a hetero atom selected from O and S, and

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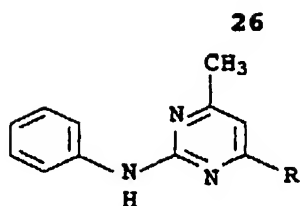
40 and [sic]

b) fungicides from the group of the dicarboximides

and/or

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c) a pyrimidine derivative of the formula III,



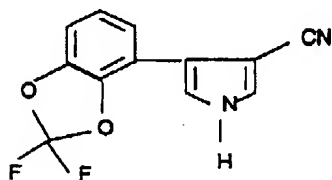
III

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in which R is methyl, propyn-1-yl or cyclopropyl,

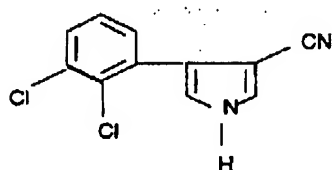
10 and/or

d) at least one active ingredient of the formula IV or V,



IV

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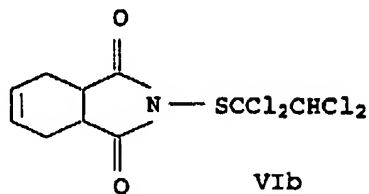
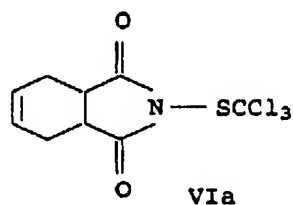


V

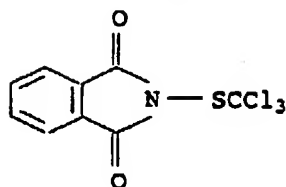
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35 and/or

e) a phthalimide derivative selected from the group consisting of the compounds VIa, VIb and VII



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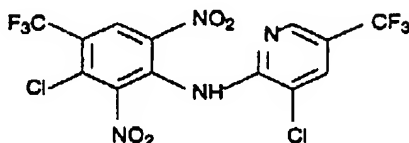
VII

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and/or

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f) a dinitroaniline of the formula VIII



VIII

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and/or

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g) an arylsulfamide of the formula IX a or IXb



IXa

25

CH₃

IXb

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in a synergistically effective amount.

2. A fungicidal mixture as claimed in claim 1, wherein in the formula I the radical A is one of the following groups:

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phenyl, pyridyl, dihydropyranyl, dihydrooxathiynyl, dihydrooxathiynyloxy, dihydrooxathiynyldioxy, furyl, thiazolyl, pyrazolyl or oxazolyl, where these groups may have 1, 2 or 3 substituents which are selected, independently of one another, from alkyl, halogen, difluoromethyl and trifluoromethyl.

40

3. A fungicidal mixture as claimed in claim 1, wherein in the formula I the radical A is one of the following groups:

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pyridin-3-yl, which may or may not be substituted in the 2-position by halogen, methyl, difluoromethyl, trifluoromethyl, methoxy, methylthio, methylsulfynyl or methylsulfonyl;

5

phenyl, which may or may not be substituted in the 2-position by methyl, trifluoromethyl, chlorine, bromine or iodine;

10

2-methyl-5,6-dihdropyran-3-yl;

2-methyl-5,6-dihydro-1,4-oxathiyn-3-yl or the 4-oxide or 4,4-dioxide thereof;

15

2-methylfuran-3-yl, which may or may not be substituted in the 4- and/or 5-position by methyl;

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thiazol-5-yl, which may or may not be substituted in the 2- and/or 4-position by methyl, chlorine, difluoromethyl or trifluoromethyl;

25

thiazol-4-yl, which may or may not be substituted in the 2- and/or 5-position by methyl, chlorine, difluoromethyl or trifluoromethyl;

30

1-methylpyrazol-4-yl, which may or may not be substituted in the 3- and/or 5-position by methyl, chlorine, difluoromethyl or trifluoromethyl; or

35

4. A fungicidal mixture as claimed in one of the preceding claims, which comprises a compound of the formula I in which R^2 is a phenyl group which may or may not be substituted by 1, 2 or 3 of the substituents mentioned in claim 1.

40

5. A fungicidal mixture as claimed in claim 4, where R^2 is a phenyl group which has one of the following substituents in the 2-position:

45

C_3 - C_6 -alkyl, C_5 - C_6 -cycloalkenyl, C_5 - C_6 -cycloalkyloxy, cycloalkenyloxy, where these groups may be substituted by 1, 2 or 3 C_1 - C_4 -alkyl groups,

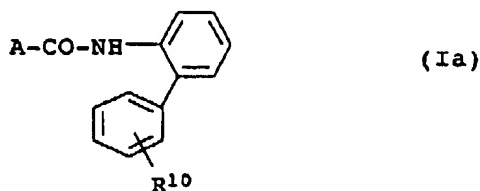
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phenyl which is substituted by from 1 to 5 halogen atoms and/or from 1 to 3 groups which are selected, independently of one another, from C₁-C₄-alkyl, C₁-C₄-haloalkyl, C₁-C₄-alkoxy, C₁-C₄-haloalkoxy, C₁-C₄-alkylthio and C₁-C₄-haloalkylthio,

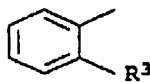
or where R² is indanyl or oxaindanyl which may or may not be substituted by 1, 2 or 3 C₁-C₄-alkyl groups.

6. A fungicidal mixture as claimed in any of claims 1 to 5, which comprises an amide compound of the formula Ia below:

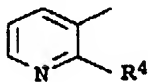


in which

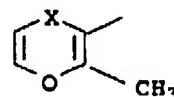
A is



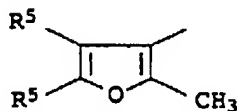
(A1)



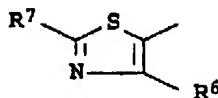
(A2)



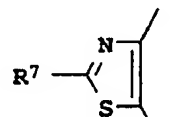
(A3)



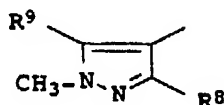
(A4)



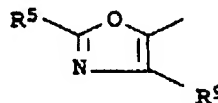
(A5)



(A6)



(A7)



(A8)

X is methylene, sulfur, sulfynyl or sulfonyl (SO₂),

R³ is methyl, difluoromethyl, trifluoromethyl, chlorine, bromine or iodine,

R⁴ is trifluoromethyl or chlorine,

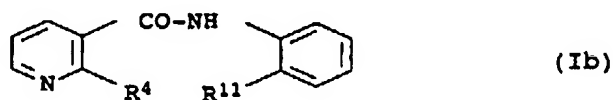
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- R^5 is hydrogen or methyl,
 R^6 is methyl, difluoromethyl, trifluoromethyl or chlorine,
 R^7 is hydrogen, methyl or chlorine,
5 R^8 is methyl, difluoromethyl or trifluoromethyl,
 R^9 is hydrogen, methyl, difluoromethyl, trifluoromethyl or chlorine,
 R^{10} is C_1 - C_4 -alkyl, C_1 - C_4 -alkoxy, C_1 - C_4 -alkylthio or halogen.

- 10 7. A fungicidal mixture as claimed in any of claims 1 to 5,
which comprises as amide compound a compound of the formula
Ib below:

15



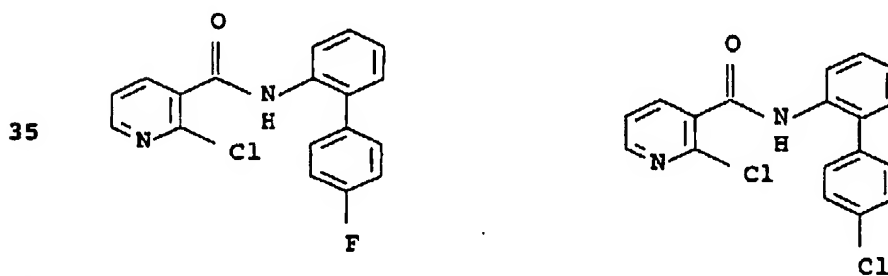
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in which

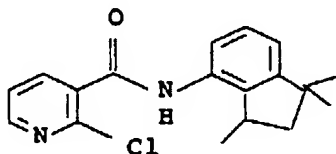
- R^4 is halogen and
25 R^{11} is phenyl which is substituted by halogen.

8. A fungicidal mixture as claimed in claim 1, which comprises
as amide compound a compound of the formulae below:

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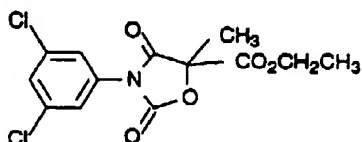


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9. A fungicidal mixture as claimed in any of the preceding claims, comprising as fungicide from the group of the dicarboximides one or more of the compounds IIa to IId

- 5 II.a: ethyl (")-3-(3,5-dichlorophenyl)-5-methyl-2,4-dioxooxazolidine-5-carboxylate [sic]

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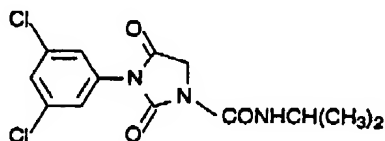


II.a

or

- 15 II.b: 3-(3,5-dichlorophenyl)-N-isopropyl-2,4-dioxoimidazolidine-1-carboxamide

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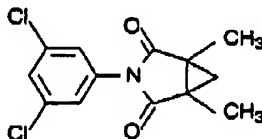


II.b

or

- 25 II.c: N-(3,5-dichlorophenyl)-1,2-dimethylcyclopropane-1,2-dicarboximide

30

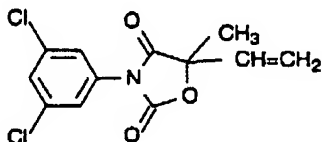


II.c

or

- 35 II.d: (RS)-3-(3,5-dichlorophenyl)-5-ethenyl-5-methyl-1,3-oxazolidine-2,4-dione

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II.d.

10. A fungicidal mixture as claimed in any of the preceding claims, which is conditioned in two parts, one part comprising the amide compound I in a solid or liquid carrier and the other part comprising one or more compounds of the formulae IIa to IXb in a solid or liquid carrier.

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11. A method for controlling harmful fungi, which comprises
treating the fungi, their habitat, or the materials, plants,
seeds, soils, areas or spaces to be protected against fungal
attack with a fungicidal mixture as claimed in any of claims
1 to 9, where the application of the active ingredients amide
compound I and one or more compounds of the formulae IIa to
IXb may be carried out simultaneously, that is either
together or separately, or in succession.

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